Pharmacovigilance Services in China

Partnering for Success in China Pharmacovigilance

If you are a pharmaceutical manufacturer, healthcare provider, or life sciences organization operating in China, access to pharmacovigilance services in China is critical to ensuring drug safety and regulatory compliance. From adverse drug reaction reporting to compliance with NMPA China pharmacovigilance guidelines, the right local pharmacovigilance partner in China shapes everything: signal detection, risk management planning, safety reporting, and successful market authorization.

Expert Pharmacovigilance Services for the China Market

China ranks as the world's second-largest pharmaceutical market, and its pharmacovigilance system has evolved significantly in recent years. With the introduction of Good Pharmacovigilance Practices (GVP) in December 2021, the National Medical Products Administration (NMPA) has standardized the responsibilities of Marketing Authorization Holders (MAHs) and clinical trial sponsors. These requirements mandate comprehensive drug safety evaluations, robust reporting systems, and continuous monitoring to reduce risks and safeguard patient health. Uniquely, China's regulatory framework extends beyond pharmaceuticals to include traditional Chinese medicine (TCM), ethnomedicine, medical devices, and cosmetics, making expert guidance vital for navigating the local drug registration and approval process.

Why Choose DDReg for Pharmacovigilance Consulting in China?

As one of the leading pharmacovigilance consulting firms , DDReg delivers end-to-end PV services in China, combining global safety expertise with local regulatory knowledge. Our experts support you in meeting all NMPA China pharmacovigilance guidelines by building robust systems for ADR reporting, aggregate safety reports, signal detection, and tailored risk management strategies. Whether you are an MAH seeking compliance or a sponsor preparing for clinical trials, DDReg ensures seamless drug safety solutions in China.

Our Pharmacovigilance & Drug Safety Services in China Include:

- + ICSR Processing & Submission
- + Risk Management Plan (RMP)
- + Aggregate Reports
- + Signal Management
- + PV Services, RPPV Functions
- + PV Audit & Compliance

Country Specific Services

- Regulatory
- Pharmacovigilance
- Labeling
- IPR Support
- Clinical Trial Regulatory
- Medical Devices regulatory services

Partner with DDReg for End-to-End Pharmacovigilance Services in China

Connect with Our Experts

Frequently Asked Questions

01 When did China's GVP come into effect?

02 Do foreign MAHs need a local PV partner in China?

03 What are the penalties for non-compliance with GVP?

04 Why choose DDReg for PV outsourcing in China?

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